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FILE COVERS 1907 - 4 Mar 2003 VOL 138 ISS 10 FILE LAST UPDATED: 3 Mar 2003 (20030303/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d que 17

L1

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L6

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L7

=> fil reg
FILE 'REGISTRY' ENTERED AT 08:45:28 ON 04 MAR 2003
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Property values tagged with IC are from the ZIC/VINITI data file

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STRUCTURE FILE UPDATES: 2 MAR 2003 HIGHEST RN 496764-40-0

DICTIONARY FILE UPDATES: 2 MAR 2003 HIGHEST RN 496764-40-0

TSCA INFORMATION NOW CURRENT THROUGH MAY 20, 2002

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

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L6 ANSWER 1 OF 7 REGISTRY COPYRIGHT 2003 ACS

RN 475304-75-7 REGISTRY

OTHER NAMES:

CN 2: PN: WO02090535 FIG: 3 claimed DNA

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RELATED SEQUENCES AVAILABLE WITH SEQLINK

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LC STN Files: CA, CAPLUS, USPATFULL

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FILE COVERS 1907 - 4 Mar 2003 VOL 138 ISS 10 FILE LAST UPDATED: 3 Mar 2003 (20030303/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

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L7 ANSWER 1 OF 9 HCAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2002:869075 HCAPLUS

DOCUMENT NUMBER: 137:365327

TITLE: The green fluorescent proteins of Renilla and Ptilosarcus and their use as reporter molecules

INVENTOR(S): Anderson, David

PATENT ASSIGNEE(S): Rigel Pharmaceuticals, Inc., USA

SOURCE: PCT Int. Appl., 130 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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WO 2002090535		A1 20021114			W	0 20	02-U	S147	66	20020509							
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	co,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	
	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,	
	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NO,	ΝZ,	OM,	PH,	
	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TR,	TT,	ΤZ,	UA,	
	UG,	US,	UZ,	VN,	YU,	ZA,	ZM,	ZW,	AM,	ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM
RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AT,	BE,	CH,	

CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG PRIORITY APPLN. INFO.:

US 2001-290287P P 20010510
US 2002-133973 A2 20020424

AB The invention relates to methods and compns. utilizing Renilla green fluorescent proteins (rGFP), and Ptilosarcus green fluorescent proteins (pGFP). In particular, the invention relates to the use of Renilla GFP or Ptilosarcus GFP proteins as reporters for cell assays, particularly intracellular assays, including methods of screening libraries, using rGFP or pGFP.

IT 475304-75-7

RL: BSU (Biological study, unclassified); BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological study); USES (Uses) (nucleotide sequence; green fluorescent proteins of Renilla and

Ptilosarcus and their use as reporter mols.)

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 2 OF 9 HCAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2001:568202 HCAPLUS

DOCUMENT NUMBER: 135:163357

TITLE: Expression of human codon modified DAF gene in mammalian cells for reducing transplant rejection

INVENTOR(S):
Miyagawa, Shuji

PATENT ASSIGNEE(S): Nippon Meat Packers, Inc., Japan SOURCE: Jpn. Kokai Tokkyo Koho, 9 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 2001211882 A2 20010807 JP 2000-22784 20000131
PRIORITY APPLN. INFO.: JP 2000-22784 20000131

AB This invention provides codon modified human complement decay-accelerating factor (DAF) gene which was expressed in transgenic mouse. The codon modification of transplant related genes is used to increase the expression of these genes in discordant transplant donor to reduce rejection reaction during transplants. The method described in this invention can be used to rejection reaction, blood coagulation and reperfusion of hypoemia.

IT 353572-08-4

RL: BOC (Biological occurrence); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study); OCCU (Occurrence); PROC (Process)

(nucleotide sequence; Expression of human codon modified DAF gene in mammalian cells for reducing rejection in transplant)

L7 ANSWER 3 OF 9 HCAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 2000:98808 HCAPLUS

DOCUMENT NUMBER: 132:146634

TITLE: Anti-angiogenesis plasmids and delivery systems and

their construction and use

INVENTOR(S): Min, Wang; Szymanski, Paul; Mehrens, Dorothy; Ralston,

Robert; Sullivan, Sean

PATENT ASSIGNEE(S): Valentis, Inc., USA SOURCE: PCT Int. Appl., 103 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent English

LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
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    WO 2000006759
                     A2
                           20000210
                                          WO 1999-US16388 19990720
    WO 2000006759
                     A3
                           20000622
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            JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK,
            MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,
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                                                          19990720
    EP 1100941
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                                        EP 1999-938769
                                                         19990720
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           AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO
                                          JP 2000-562541
                                                          19990720
    JP 2002524036
                     T2 20020806
PRIORITY APPLN. INFO.:
                                       US 1998-94375P P 19980727
                                       WO 1999-US16388 W 19990720
```

AB The present invention relates to gene delivery and gene therapy, and provides novel nucleic acid constructs for expression of anti-angiogenic agents in a mammal, formulations for delivery that incorporate a nucleic acid construct for expression, and methods for prepg. and using such constructs and formulations. In particular, this invention relates to plasmid constructs for delivery of therapeutic anti-angiogenic encoding nucleic acids to cells in order to modulate tumor activity, methods of using those constructs (including combination therapy with other agents, such as cytokines, preferably interleukin-12), as well as methods for prepg. such constructs. Plasmid vectors are constructed comprising synthetic genes having optimal codon usage for endostatin and angiostatin expression, under the control of a promoter specific for expression in endothelial cells (e.g., the enhancer of cytomegalovirus for human endothelin-1) and the growth hormone 3'-untranslated region with a deleted Alu repeat. A polymeric gene delivery system uses polyvinyl pyrrolidone to increase protein expression by protecting plasmid DNA fron nucleases and controlling the dispersion and retention of plasmid DNA in injected tissues. The plasmid delivery system also comprises a cationic lipid (DOPTMA), neutral lipid (cholesterol), and an isotonic carbohydrate (lactose) soln.

258258-38-7 TΤ

RL: BUU (Biological use, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(nucleotide sequence; anti-angiogenesis plasmids and delivery systems and their construction and use)

ANSWER 4 OF 9 HCAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1999:614138 HCAPLUS

DOCUMENT NUMBER: 131:238811

Expression vectors for interferon .alpha. genes for TITLE:

use in gene therapy

Nordstrom, Jeff; Pericle, Federica; Rolland, Allain; INVENTOR(S):

Ralston, Robert

PATENT ASSIGNEE(S):

Genemedicine, Inc., USA

SOURCE:

PCT Int. Appl., 137 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent English

LANGUAGE:

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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APPLICATION NO. DATE
                 KIND DATE
    PATENT NO.
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    WO 9947678
                           19990923
                                         WO 1999-US5394 19990312
                     A2
                     A3 19991209
    WO 9947678
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            DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS,
            JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK,
            MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD,
            RU, TJ, TM
        RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK,
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            CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
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                                        CA 1999-2323604 19990312
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    AU 9930003
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                      A1
                                         EP 1999-911340
                                                         19990312
    EP 1064383
                          20010103
                     A2
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, FI
                                          JP 2000-536861 19990312
    JP 2002506647
                     Т2
                           20020305
                                       US 1998-78654P P 19980319
PRIORITY APPLN. INFO.:
                                       WO 1999-US5394 W 19990312
```

Plasmid expression constructs for mammalian interferon .alpha. genes that can be used in gene therapy and methods for delivering them are described. These constructs can be used in the treatment of cancer, including combination therapy with other agents, such as cytokines, preferably IL-12. Constructs using a cytomegalovirus promoter, a synthetic 5'-intron and the 3'-UTR of the human growth hormone gene are described. The interferon .alpha. genes may have its codon usage altered to maximize or increase expression. Direct administration of a mouse interferon .alpha.4 gene to renal cell carcinomas and mammary cell adenocarcinomas was shown to inhibit tumor growth, and bring about complete complete regression in some cases. The treatment also induced long-lasting immunity to secondary tumor challenges.

207241-26-7 TΤ

> RL: BSU (Biological study, unclassified); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses) (nucleotide sequence; expression vectors for interferon .alpha. genes for use in gene therapy)

ANSWER 5 OF 9 HCAPLUS COPYRIGHT 2003 ACS 1998:550439 HCAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER: 129:185076

TITLE: IL-2 gene expression, liposome delivery system, and

uses

INVENTOR(S): Ralston, Robert; Muller, Susanne; Mumper, Russ;

Munger, William; Bruno, Maria

PATENT ASSIGNEE(S):

Genemedicine, Inc., USA

SOURCE: PCT Int. Appl., 77 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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PATENT NO.
                 KIND DATE
                                         APPLICATION NO. DATE
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    WO 9834952
                          19980813
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            NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
            UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI,
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            GA, GN, ML, MR, NE, SN, TD, TG
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    EP 975780
                                       EP 1998-904943 19980209
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                          20000202
           AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, FI
    JP 2001511647
                    Т2
                          20010814
                                         JP 1998-534889
                                                        19980209
PRIORITY APPLN. INFO.:
                                      US 1997-39709P P 19970210
                                      US 1998-12366
                                                     A 19980123
                                      WO 1998-US2221
                                                    W 19980209
```

AB Plasmid expression systems for delivery of DNA coding sequences in liposomes to a mammal are described which provide expression of human IL-2 for therapy of cancers and other diseases. Also described are particular lipid/DNA delivery systems having advantageous characteristics of size, charge ratio, and proportion of supercoiled DNA, and methods of prepg. and using such delivery systems for treatment.

L7 ANSWER 6 OF 9 HCAPLUS COPYRIGHT 2003 ACS ACCESSION NUMBER: 1998:398430 HCAPLUS

DOCUMENT NUMBER: 129:64088

TITLE: Insulin-like growth factor I (IGF-I) expression vector

for gene therapy

INVENTOR(S): Coleman, Michael; Schwartz, Robert; Demayo, Francesco

J.

PATENT ASSIGNEE(S): GeneMedicine, Inc., USA; Baylor College of Medicine

SOURCE: PCT Int. Appl., 116 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

and uses)

PATENT INFORMATION:

PATENT NO.			KIND D		DATE			APPLICATION NO.					DATE						
								_											
	WO 9824922			A1 19980611				W	0 19	97-U	S218	52	19971201						
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			DK,	EE,	ES,	FI,	GB,	GE,	GH,	ΗU,	ID,	IL,	IS,	JP,	ΚE,	KG,	ΚP,	KR,	
			KZ.	LC,	LK.	LR.	LS,	LT.	LU.	LV,	MD,	MG,	MK,	MN,	MW,	MX,	NO,	NZ,	

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RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA,
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     AU 9853662
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                             19980629
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     EP 943003
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                             19990922
                                                                19971201
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
              IE, FI
     JP 2001505435
                        T2
                              20010424
                                              JP 1998-525696
                                                                19971201
     US 2003018984
                                              US 2001-861101
                        Α1
                              20030123
                                                                20010518
                                           US 1996-31539P P 19961202
PRIORITY APPLN. INFO.:
                                                            A 19971119
                                          US 1997-974572
                                          WO 1997-US21852 W 19971201
AB
     This invention relates to gene delivery and expression, including gene
     therapy, by using vectors which encode stable mRNA and methods of using
     such vectors. In particular, this invention relates to vectors which
     establish controlled expression of recombinant IGF-I genes within tissues
     at certain levels. The vector includes a 5' flanking region which
     includes necessary sequences for expression of a nucleic acid cassette, a
     3' flanking region including a 3' UTR and/or 3' NCR, and a linker which
     connects the 5' flanking region to a nucleic acid sequence. The linker
     has a position for inserting a nucleic acid cassette. The linker does not
     contain the coding sequence of a gene that the linker is naturally assocd.
     with. The 3' flanking region is 3' to the position for inserting the
     nucleic acid cassette. The expression vectors of the present invention
     can also be regulated by a regulatory system and/or constructed with a
     coating. Expression plasmids contg. chicken skeletal muscle .alpha.-actin
     gene promoter and first intron, human IGF-I cDNA and human growth hormone
     gene 3'-UTR and poly(A) signal were prepd. Replacement of skeletal muscle
     .alpha.-actin gene 3'-UTR of prior art plasmids with the 3'-UTR of human
     growth hormone (as above) resulted in increased delivery of IGF-I from
     skeletal muscle to systemic circulation in both transgenic animal (mouse)
     and non-viral gene therapy paradigms. The vector was nontoxic in
     safety/toxicol. studies in dogs. Use of the vector in treatment of
     diabetes (in rats) and (calf) muscle disuse atrophy was demonstrated.
     209056-07-5
TT
     RL: BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological
     study); USES (Uses)
        (nucleotide sequence; insulin-like growth factor I expression vector
        for gene therapy)
REFERENCE COUNT:
                                 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS
                                 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 7 OF 9 HCAPLUS COPYRIGHT 2003 ACS
                          1998:268632 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                          129:50507
TITLE:
                          Polycistronic expression constructs for gene therapy
                          using interleukin 12 genes and their delivery by
                          liposomes
                          Nordstrom, Jeff; Freimark, Bruce; Deshpande, Deepa
INVENTOR(S):
                          Genemedicine, Inc., USA; Nordstrom, Jeff; Freimark,
PATENT ASSIGNEE(S):
                          Bruce; Deshpande, Deepa
                          PCT Int. Appl., 105 pp.
SOURCE:
                          CODEN: PIXXD2
DOCUMENT TYPE:
                          Patent
LANGUAGE:
                          English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
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APPLICATION NO. DATE
     PATENT NO. KIND DATE
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                         A2
     WO 9817814
                                               WO 1997-US18832 19971010
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                        A3 19980827
     WO 9817814
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RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, CN, MI, MB, NE, SM, TD, TG
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     JP 2001503258
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     US 2002119940 A1
                                                US 2001-754014 20010103
                               20020829
                                             US 1996-28687P P 19961018
PRIORITY APPLN. INFO.:
                                             US 1997-948958 B1 19971010
                                             WO 1997-US18832 W 19971010
     Plasmid expression vectors for delivery of genes for a heterooligomeric
AΒ
     proteins that ensure coordinated expression of the genes are described for
     use in gene therapy. In particular, these vectors use synthetic introns
     and 5'- and 3'-untranslated regions to control expression. These
     sequences are designed to have good fits with consensus sequences for
     important functional sequences such as Kozak boxes and intron cleavage and
     splice sites. Also described are particular lipid/DNA delivery systems
     having advantageous characteristics of size, charge ratio, and proportion
     of supercoiled DNA, and methods of prepg. and using such delivery systems
     for treatment or as immunization adjuvants. Optimization expts. are
     described. Studies with a guinea pig model of asthma showed that
     liposomes of the expression cassette introduced into the bronchi could be
     used to reduce the inflammatory response.
     207241-26-7
IT
     RL: BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological
     study); USES (Uses)
         (nucleotide sequence; polycistronic expression constructs for gene
         therapy using interleukin 12 genes and their delivery by liposomes)
     ANSWER 8 OF 9 HCAPLUS COPYRIGHT 2003 ACS
                      1998:65983 HCAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                           128:150393
                           Purification and recombinant production of human
TITLE:
                           telomerase subunits and their applications for drug
                           screening and therapy
                           Cao, Zhaodan
INVENTOR(S):
PATENT ASSIGNEE(S):
                           Tularik, Inc., USA
                            PCT Int. Appl., 33 pp.
SOURCE:
                           CODEN: PIXXD2
DOCUMENT TYPE:
                            Patent
                            English
LANGUAGE:
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
     PATENT NO. KIND DATE
                                              APPLICATION NO. DATE
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     WO 9801543 A1 19980115
                                                WO 1997-US12297 19970708
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                                         US 1996-676967 19960708
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                                                              19970708
     US 5888747
                                            US 1998-72270
                       Α
                             19990330
                                                              19980504
PRIORITY APPLN. INFO.:
                                         US 1996-676967
                                                              19960708
                                         WO 1997-US12297
                                                              19970708
     The invention provides methods and compns. relating to a human telomerase
AΒ
     and related nucleic acids, including 4 distinct human telomerase subunit
     proteins called p140, p105, p48 and p43 having human telomerase-specific
     activity. Human telomerase p105 subunit cDNA contains an open reading frame encoding 759 amino acids. The proteins may be produced
     recombinantly from transformed host cells from the disclosed telomerase
     encoding nucleic acids or purified from human cells. Also included are human telomerase RNA components, as well as specific, functional derivs.
     thereof. The invention provides isolated telomerase hybridization probes
     and primers capable of specifically hybridizing with the disclosed
     telomerase gene, telomerase-specific binding agents such as specific
     antibodies, and methods of making and using the subject compns. in
     diagnosis, therapy and in the biopharmaceutical industry.
ΙT
     202485-81-2
     RL: BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological
     study); USES (Uses)
        (nucleotide sequence; purifn. and recombinant prodn. of human
        telomerase subunits and their applications for drug screening and
        therapy)
     ANSWER 9 OF 9 HCAPLUS COPYRIGHT 2003 ACS
                         1998:65982 HCAPLUS
ACCESSION NUMBER:
                         128:151114
DOCUMENT NUMBER:
                         Purification and recombinant production of human
TITLE:
                         telomerase subunits and their applications for drug
                         screening and therapy
INVENTOR(S):
                         Collins, Kathleen
                         Regents of the University of California, USA
PATENT ASSIGNEE(S):
                         PCT Int. Appl., 33 pp.
SOURCE:
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
                         English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
                                    APPLICATION NO. DATE
     PATENT NO.
                  KIND DATE
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                     A1 19980115 WO 1997-US12296 19970708
     WO 9801542
         W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
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             LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL,
             PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ,
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                       Α
PRIORITY APPLN. INFO.:
                                         US 1996-676974
                                                             19960708
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WO 1997-US12296

19970708

and related nucleic acids, including 4 distinct human telomerase subunit proteins called p140, p105, p48 and p43 having human telomerase-specific activity. Human telomerase p105 subunit cDNA contains an open reading frame encoding 759 amino acids. The proteins may be produced recombinantly from transformed host cells from the disclosed telomerase encoding nucleic acids or purified from human cells. Also included are human telomerase RNA components, as well as specific, functional derivs. thereof. The invention provides isolated telomerase hybridization probes and primers capable of specifically hybridizing with the disclosed telomerase gene, telomerase-specific binding agents such as specific antibodies, and methods of making and using the subject compns. in diagnosis, therapy and in the biopharmaceutical industry.

IT 202485-81-2

RL: BUU (Biological use, unclassified); PRP (Properties); BIOL (Biological study); USES (Uses)

(nucleotide sequence; purifn. and recombinant prodn. of human telomerase subunits and their applications for drug screening and therapy)